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OM protein - protein search, using sw model

Run on: December 24, 2002, 22:08:57 ; Search time 68 Seconds
(without alignments)
811.261 Million cell updates/sec

Title: US-09-708-724a-2

Perfect score: 2187

Sequence: 1 MGPSVSVVLCVCGHAGL.....LLAVTRGLELRIISKRRAE 414

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 908470 seqs, 133250620 residues

Total number of hits satisfying chosen parameters: 908470

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listed first 45 summaries

Database : A_Geneseq_101002.*

- 1: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1980.DAT.*
- 2: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1981.DAT.*
- 3: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1982.DAT.*
- 4: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1983.DAT.*
- 5: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1984.DAT.*
- 6: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1985.DAT.*
- 7: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1986.DAT.*
- 8: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1987.DAT.*
- 9: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1988.DAT.*
- 10: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1989.DAT.*
- 11: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1990.DAT.*
- 12: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1991.DAT.*
- 13: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1992.DAT.*
- 14: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1993.DAT.*
- 15: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1994.DAT.*
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- 18: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1997.DAT.*
- 19: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1998.DAT.*
- 20: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA1999.DAT.*
- 21: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA2000.DAT.*
- 22: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA2001.DAT.*
- 23: /SIDS2/gcgdata/geneseq/geneseq-emb1/AA2002.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	ID	Description
1	259.5	11.9	334	22 ABG04612	Novel human diago
2	258	11.7	133	22 ABA41145	Peptide #8651 enco
3	256	11.7	133	22 AAMG2001	Human brain expres
4	256	11.7	133	22 AAM74805	Human bone marrow
5	256	11.7	133	22 AAM34921	Peptide #8958 enco
6	256	11.7	133	23 ABG44596	Human peptide enco
7	254	11.6	122	22 ABG04611	Novel human diago
8	130	5.9	1178	22 ABB68342	Drosophila melanog
9	129.5	5.9	641	22 AAM38978	Human polypeptide
10	129.5	5.9	698	22 AAM38977	Human polypeptide

11	129.5	5.9	698	22 AAM39032	Human polypeptide
12	129.5	5.9	705	22 AAM39135	Human protein sequ
13	129.5	5.9	706	22 AAM38979	Human polypeptide
14	129.5	5.9	1837	21 AAY85564	Human homologue of
15	128.5	5.9	828	22 ABB65774	Drosophila melanog
16	128	5.9	1593	23 AAM48935	Murine MEKK1-2. M
17	125.5	5.7	595	22 AAE01114	Human gene 1 encod
18	125.5	5.7	595	23 ABG64591	Human albumin fusi
19	125.5	5.7	1560	21 AAB18792	The human ribosome
20	125	5.7	1328	22 AAM78519	Human protein SEQ
21	125	5.7	1331	22 AAM79503	Human protein SEQ
22	123.5	5.6	2639	22 ABB15016	Novel human diago
23	120.5	5.5	608	22 AAB20164	Human protein asso
24	118	5.4	482	22 AAB31195	Amino acid sequenc
25	118	5.4	482	22 AAY97586	Human secreted pro
26	118	5.4	482	22 AAB65298	Human PRO170 prot
27	118	5.4	482	22 AAB27225	Human EXMAD-3 SEQ
28	118	5.4	538	22 AAM38985	Human polypeptide
29	118	5.4	538	22 AAE06598	Human protein havi
30	117	5.3	905	18 AAW31186	Human p160 polypep
31	116	5.3	561	22 AAM40711	Human polypeptide
32	115	5.3	503	22 AAB20165	Human protein asso
33	114	5.2	2161	22 AAM78959	Human protein SEQ
34	114	5.2	2189	22 AAM79943	Human protein SEQ
35	114	5.2	2523	22 AAU03503	Human protein kina
36	113.5	5.2	401	21 AAB41664	Human ORFX ORF1428
37	113.5	5.2	425	22 AAB55309	Protonibacterium
38	113.5	5.2	1721	19 AAM48299	Cryptosporidium pa
39	113	5.2	1145	22 AAU04895	Micromonospora eve
40	113	5.2	2429	23 AAE21713	Human PKIN-8 prote
41	112.5	5.1	584	23 AAE23791	Human BCAS1 (brea
42	112.5	5.1	594	22 ABB16928	Novel human diago
43	112.5	5.1	914	22 ABB69998	Drosophila melanog
44	112.5	5.1	1331	22 ABB28241	Novel human diago
45	112	5.1	712	23 ABB61500	Human NF-KB activa

*ALIGNMENTS

RESULT 1

ABG04612
ID ABG04612 standard; Protein; 334 AA.

XX AC ABG04612;

XX DT 13-FEB-2002 (first entry)

XX DE Novel human diagnostic protein #4603.

XX KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
XX KW food supplement; medical imaging; diagnostic; genetic disorder.

XX OS Homo sapiens.

XX PN WO200175067-A2.

XX PD 11-OCT-2001.

XX PF 30-MAR-2001; 2001WO-US08631.

XX PR 31-MAR-2000; 2000US-0540217.

XX PR 23-AUG-2000; 2000US-0649167.

XX XX (HYSE-) HYSEQ INC.

XX PI Drmanac RT, Liu C, Tang YT;

XX DR WPI; 2001-639362/73.

XX DR N-PSDB; AAS68799.

XX PT New isolated polynucleotide and encoded polypeptides, useful in
XX PT diagnostics, forensics, gene mapping, identification of mutations

QY 296 TIAGEPLGHCHTFTIS 310
 DB 114 ATAGEPSGCAFPIS 128

RESULT 6
 ABG04596
 ID ABG04596 standard; Peptide; 133 AA.
 AC ABG04596;
 XX
 DT 19-AUG-2002 (first entry)
 XX
 DE Human peptide encoded by genome-derived single exon probe SEQ ID 34261.
 KW Human: single exon probe; asthma; lung cancer; COPD; ILD;
 KW chronic obstructive pulmonary disease; interstitial lung disease;
 KW familial idiopathic pulmonary fibrosis; neurofibromatosis;
 KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
 KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
 KW pulmonary histiocytosis; lymphangioleiomyomatosis; Karagener syndrome;
 KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
 KW primary ciliary dyskinesia; pulmonary hypertension;
 KW hyaline membrane disease.
 XX
 OS Homo sapiens.
 XX
 PN WO200186003-A2.
 XX
 XX 15-NOV-2001.
 PD
 XX
 PF 30-JAN-2001; 2001WO-US00665.
 XX
 PR 04-FEB-2000; 2000US-180312P.
 PR 26-MAY-2000; 2000US-207456P.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0632366.
 PR 21-SEP-2000; 2000US-234687P.
 PR 27-SEP-2000; 2000US-236359P.
 PR 04-OCT-2000; 2000GB-0024263.
 XX
 XX (MOLE-) MOLECULAR DYNAMICS INC.
 PA
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 XX WPI; 2002-114183/15.
 DR
 XX Spatially-addressable set of single exon nucleic acid probes, used to
 PT measure gene expression in human lung samples -
 XX
 XX Claim 27; SEQ ID No 34261; 634pp; English.
 PS
 CC The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human lung comprising single exon nucleic acid probes having one of
 CC 12614 nucleic acid sequences mentioned in the specification, or their
 CC complements or the 12387 open reading frames derived from the 12614
 CC probes. Also included are a microarray comprising the novel set of
 CC probes; the novel set of probes which hybridise at high stringency to a
 CC nucleic acid expressed in the human lung; measuring gene expression in a
 CC sample derived from human lung, comprising (a) contacting the array with
 CC a collection of detectably labeled nucleic acids derived from human lung
 CC mRNA, and (b) measuring the label detectably bound to each probe of
 CC the array; identifying exons in a eukaryotic genome, comprising
 CC (a) algorithmically predicting at least one exon from genomic sequences
 CC of the eukaryote; and (b) detecting specific hybridisation of detectably
 CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
 CC having a fragment identical to the predicted exon, the probe is included
 CC in the above mentioned microarray; assigning exons to a single gene,
 CC comprising (a) identifying exons from genomic sequence by the method
 CC above and (b) measuring the expression of each of the exons in several
 CC tissues and/or cell types using hybridisation to a single exon

CC microarrays having a probe with the exon, where a common pattern of
 CC expression of the exons in the tissues and/or cell types indicates that
 CC the exons should be assigned to a single gene; a peptide comprising one
 CC of 12011 sequences, mentioned in the specification, or encoded by the
 CC probes/open reading frames (ORF). The probes are used for gene
 CC expression analysis, and for identifying exons in a gene, particularly
 CC using human lung derived mRNA and for the study of lung diseases
 CC such as asthma, lung cancer, chronic obstructive pulmonary disease
 CC (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
 CC fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
 CC Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
 CC haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis,
 CC pulmonary alveolar proteinosis, Karagener syndrome, fibrocystic
 CC pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
 CC and hyaline membrane disease. The present sequence is a peptide/protein
 CC encoded by a single exon probe of the invention.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 133 AA;
 Query Match 11.7%; Score 256; DB 23; Length 133;
 Best Local Similarity 68.0%; Pred. No. 2.7e-13;
 Matches 51; Conservative 6; Mismatches 18; Indels 0; Gaps 0;
 QY 236 SESQILKESFVPTTPKENNKQEREDENRLLPPPPVAETPPSPVSTETETPLQIRPTA 295
 DB 54 AESKMLKESVPPPTASIKKQEREDKNWPIILPPVAETSPVPPSPVAGIETPIQILKSA 113
 QY 296 TIAGEPLGHCHTFTIS 310
 DB 114 ATAGEPSGCAFPIS 128

RESULT 7
 ABG04611
 ID ABG04611 standard; Protein; 122 AA.
 AC ABG04611;
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE Novel human diagnostic protein #4602.
 KW Human: chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder.
 XX
 OS Homo sapiens.
 XX
 PN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-US08631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HYSE-) HYSEQ INC.
 PI Drmanac RT, Liu C, Tang YT;
 XX
 XX WPI; 2001-639362/73.
 DR N-PSDB; AAS68798.
 XX
 PT New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 PS Claim 20; SEQ ID No 34970; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. ABG00010-ABG30377 represent novel human
CC diagnostic amino acid sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 122 AA;

Query Match 11.6%; Score 254; DB 22; Length 122;
Best Local Similarity 59.1%; Pred. No. 3.5e-13;
Matches 52; Conservative 7; Mismatches 21; Indels 8; Gaps 1;

QY 236 SESQILKESFVPTTPKNNKQERDENWRLPPPPVATPVPSPSVTEIPLQRIPTA 295
DB 33 AESNLKESVVPPTAPTENKQVREDKNWIPPPPIAETSVLPSPVABETPKOTLCSA 92

QY 296 TIAGEPLGCHCTFTIS-----PAFVH 315
DB 93 AIAGEPLGCTFPISVRPSNNPQQFIH 120

RESULT 8
ABB68342

ID ABB68342 standard; Protein; 1178 AA.

XX ABB68342;

DT 26-MAR-2002 (first entry)

DE Drosophila melanogaster polypeptide SEQ ID NO 31818.

XX Drosophila; developmental biology; cell signalling; insecticide;
KW pharmaceutical.

XX Drosophila melanogaster.

XX WO200171042-A2.

PN 27-SEP-2001.

XX 23-MAR-2001; 2001WO-US09231.

XX 23-MAR-2000; 2000US-191637P.

PR 11-JUL-2000; 2000US-0614150.

XX (PEKE) PE CORP NY.

XX Venter JC, Adams M, Li PWD, Myers EW;

XX WPI; 2001-656860/75.

DR N-PSDB; ABL12445.

XX New isolated nucleic acid detection reagent for detecting 1000 or more
PT genes from Drosophila and for elucidating cell signalling and cell-cell
PT interactions -

XX

PS Disclosure; SEQ ID NO 31818; 21pp + Sequence Listing; English.

XX The invention relates to an isolated nucleic acid detection reagent
CC capable of detecting 1000 or more genes from Drosophila. The invention is
CC useful in developmental biology and in elucidating cell signalling and
CC cell-cell interactions in higher eukaryotes for the development of
CC insecticides, therapeutics and pharmaceutical drugs. The invention
CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
CC sequences (ABL01840-ABL16175) and the encoded proteins
CC (ABB57737-ABB72072).

CC The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 1178 AA;

Query Match 5.9%; Score 130; DB 22; Length 1178;
Best Local Similarity 19.8%; Pred. No. 0.12;
Matches 99; Conservative 59; Mismatches 180; Indels 162; Gaps 19;

QY 49 QNPTLPSPSHRPPGNAASVVTGGDCHLPTTEEFGLV-----QSMKCDTVRIKGVLIQ 101
DB 416 QKPQVTCVTLVPPPOESPVSSAAETLEKRPQVGAISTHESPRQSLSSPTDSVDSAKS 475

QY 102 GPTTAPPLMTS-----EGNVTAEDEEIAFAV-----YAV 132
DB 476 TPSASPKPQTQVPMQPLLQMRPHLQANLSAQ-TPVTVPARVVRMQQAQIIPQRLPV 534

QY 133 AASASAEAAWHRLV-----LLSGQIHEPI-----GSGG 161
DB 535 ASSAAMATMTRSIVTSTAGTSTITGR---PVATTLNNSNPTVAQLQSMANMAGGGG 591

QY 162 NIINTNG-----GRSC-----ONPA-----LPSDQ 183
DB 592 QLIMTSSGQLLVIPTPSKQTTQHHRRPGPGQGVIIQQQPAELHPQGGYIVSQSPVA 651

QY 184 SPNGNATTSVTRDN-----YHLLTEEFVWSQSMKWHSONK-----SGGSVP 226
DB 652 AASSSSSTVILNSGGAKLLHHQIITSQAGQINQATSGSGNQPTVLLNPLNGYIV 711

QY 227 VRGP-TQEPCESEQILKESFVPP-----TTPKNNKQERDENWRLPPPPVAE-----T 274
DB 712 QQQPQTQSPQAEQILAMPQPPPAQTLIISPDTKRRARKKSSVCHTPPPSPGSPAKIIS 771

QY 275 PVPSPVTEIETPLQRIPTATIAEGLGCHCTFTISFAFVHSLNKR----- 322
DB 772 FQISPSINQAPALLHQAAAAAQAAPQOQFQLSPGIQIVVKNPQPPQPTQQQLLL 831

QY 323 -----RQLEILLREVEWPGRHMAATCCKLQVEGQDRT-MSLAAAPVREAPPPPTGASS 375
DB 832 QNGQILQOVNLIGQQLLMPAGLVMPGPDATLLIQNMPATSLMTPOGPVMLRTPSPQNKPS 891

QY 376 --EPSVPA---LPGADPQRS 390

DB 892 FISPAGGQQYLVGANGQLS 911

RESULT 9
AAM38978

ID AAM38978 standard; Protein; 641 AA.

XX AAM38978;

AC AAM38978;

DT 22-OCT-2001 (first entry)

XX Human polypeptide SEQ ID NO 2123.

XX Human; nootropic; immunosuppressant; cytostatic; gene therapy; cancer;
KW peripheral nervous system; neuropathy; central nervous system; CNS;
KW Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
KW amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
KW chemokinetic; thrombolytic; drug screening; arthritis; inflammation;
KW leukaemia.

DE Human protein sequence SEQ ID NO:14398.
 KW Human; primer; detection; diagnosis; antisense therapy; gene therapy.
 OS Homo sapiens.
 PN EP1074617-A2.
 XX 07-FEB-2001.
 PD 28-JUL-2000; 2000EP-0116126.
 PF 29-JUL-1999; 99JP-0248036.
 PR 17-AUG-1999; 99JP-0300253.
 PR 11-JAN-2000; 2000JP-0118776.
 PR 02-MAY-2000; 2000JP-0183767.
 PR 09-JUN-2000; 2000JP-0241899.
 XX (HELI-) HELIX RES INST.
 PA Ota T, Isoqai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
 PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
 XX WPI: 2001-318749/34.
 DR Primer sets for synthesizing polynucleotides, particularly the 5602
 PT full-length cDNAs defined in the specification, and for the detection
 PT and/or diagnosis of the abnormality of the proteins encoded by the
 PT full-length cDNAs -
 XX Claim 8; SEQ ID 14398; 2537pp + CD ROM; English.
 CC The present invention describes primer sets for synthesizing 5602
 CC full-length cDNAs defined in the specification. Where a primer set
 CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
 CC to the complementary strand of a polynucleotide which comprises one of
 CC the 5602 nucleotide sequences defined in the specification, where the
 CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination
 CC of an oligonucleotide comprising a sequence complementary to the
 CC complementary strand of a polynucleotide which comprises a 5'-end
 CC sequence and an oligonucleotide comprising a sequence complementary to a
 CC polynucleotide which comprises a 3'-end sequence, where the
 CC oligonucleotide comprises at least 15 nucleotides and the combination of
 CC the 5'-end sequence/3'-end sequence is selected from those defined in
 CC the specification. The primer sets can be used in antisense therapy and
 CC in gene therapy. The primers are useful for synthesizing polynucleotides,
 CC particularly full-length cDNAs. The primers are also useful for the
 CC detection and/or diagnosis of the abnormality of the proteins encoded by
 CC the full-length cDNAs. The primers allow obtaining of the full-length
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to
 CC AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
 CC represent oligonucleotides, all of which are used in the exemplification
 CC of the present invention.
 XX Sequence 705 AA;
 SQ Query Match 5.9%; Score 129.5; DB 22; Length 705;
 Best Local Similarity 21.9%; Pred. No. 0.066;
 Matches 89; Conservative 45; Mismatches 165; Indels 107; Gaps 17;
 QY 41 GAQNLMTCONP-----TLPSVSHRSP-----GNAASVTGGDCHLPTFEFGVLVQSM 89
 DB 90 GRDLSDAKKPPSGIARPTSGSGFYKKPPATGATVMTGTGS-----ATLSKIQ 140
 QY 90 KCDTVRIKVGLOPPTAPPLTSEGNTVAEDTEAIRFAVVAASAAEAHWHRLVLL 149
 DB 141 KSGGIPVKPV-----NCRKTLSDVSNSEAPGFLAPGARSNIQ-----YRSL---- 181
 QY 150 SQGHEPIGSGGNIINTNKGRCQNPALPSPDQSPGNGATTSVTDNHYLLTEEEFGVW 209
 DB 182 ----PRPAKSSMSVT---GGRGGRPRVSSSIDPS-----LLSTQOGGLT 219

QY 210 SOSMKWHSQKSGGVPVGRPTQPCSESIKESFV-----PPTTPKNN 255
 DB 220 PSLKEPTKVASGRTP--APVQTDREKAKAKAVALDSNISLKSIGSPSTPKNQ 277
 QY 256 KOEREDENRLLPPPV--AETPVSPSVTEIE-----TPLORIP---RT 294
 DB 278 SHPTATKLAELPPTPLRATAKSFVKPPPSLANLDKVNNSLDLPSSDTHASKVPDLHAT 337
 QY 295 ATIAGEPLGCHCTTISPAFVHSLNKRKRLLELLREVEWPGRHMAATCCKL--QVEGQ 352
 DB 338 SSASGGPLPSC--FTSPAPILNINSASFQGLELMSGFSVPKRTMYPKLSGLHRSMSL 396
 QY 353 DRTMSLAAA-----PVREAPPPTGASSESPVPALPGADPQSAEL 393
 DB 397 QMPMSLPSAFTSPV-PTPPAPPAAPTEETEELTWSSGSPRAGL 441
 RESULT 13
 AAM38979
 ID AAM38979 standard; Protein; 706 AA.
 AC AAM38979;
 XX
 DT 22-OCT-2001 (first entry)
 XX Human polypeptide SEQ ID NO 2124.
 DE Human; nontropic; immunosuppressant; cytostatic; gene therapy; cancer;
 KW peripheral nervous system; neuropathy; central nervous system; CNS;
 KW Alzheimer's; Parkinson's disease; Huntington's disease; haemostatic;
 KW amyotrophic lateral sclerosis; Shy-Drager Syndrome; chemotactic;
 KW chemokinetic; thrombolytic; drug screening; arthritis; inflammation;
 KW leukaemia.
 XX Homo sapiens.
 OS WO200153312-A1.
 PN 26-JUL-2001.
 PD 26-DEC-2000; 2000WO-US34263.
 PF 21-JAN-2000; 2000US-0488725.
 PR 25-APR-2000; 2000US-0552317.
 PR 09-JUL-2000; 2000US-0598042.
 PR 19-JUL-2000; 2000US-0620312.
 PR 03-AUG-2000; 2000US-0653450.
 PR 14-SEP-2000; 2000US-0662191.
 PR 19-OCT-2000; 2000US-0693036.
 PR 29-NOV-2000; 2000US-0727344.
 XX (HYSE-) HYSEQ INC.
 PA Tang YT, Liu C, Asundi V, Chen R, Ma Y, Qian XB, Ren F, Wang D;
 PI Wang J, Wang Z, Wehrman T, Xu C, Xue AJ, Yang Y, Zhang J;
 PI Zhao QA, Zhou P, Goodrich R, Drmanac RT;
 XX WPI: 2001-442253/47.
 DR N-FSDB; AA158135.
 DR Novel nucleic acids and polypeptides, useful for treating disorders
 PT such as central nervous system injuries -
 PS Example 4; SEQ ID NO 2124; 10078pp; English.
 XX The invention relates to human nucleic acids (AA157798-AA161369) and
 CC the encoded polypeptides (AAM38642-AA42213) with nontropic,
 CC immunosuppressant and cytostatic activity. The polynucleotides are useful
 CC in gene therapy. A composition containing a polypeptide or polynucleotide
 CC of the invention may be used to treat diseases of the peripheral nervous
 CC system, such as peripheral nervous injuries, peripheral neuropathy and
 CC localised neuropathies and central nervous system diseases, such as
 CC Alzheimer's, Parkinson's disease, Huntington's disease, amyotrophic

CC lateral sclerosis, and Shy-Drager Syndrome. Other uses include the
 CC utilisation of the activities such as: Immune system suppression,
 CC Activin/inhibin activity, chemotactic/chemokinetic activity, haemostatic
 CC and thrombolytic activity, cancer diagnosis and therapy, drug screening,
 CC assays for receptor activity, arthritis and inflammation, leukemias and
 CC C.N.S disorders.
 CC Note: The sequence data for this patent did not form part of the printed
 CC specification.
 XX
 SQ

Sequence 706 AA;

Query Match 5.9%; Score 129.5; DB 22; Length 706;
 Best Local Similarity 21.9%; Pred. No. 0.066;
 Matches 89; Conservative 45; Mismatches 165; Indels 107; Gaps 17;

QY 41 GAQNLMTQNP-----TLPSVSHRSP---GNAAVSVTGDCCHLTTEEEFGVLVQSM 89
 DB 90 GDRLSDAKKPPSGTARSTSGSGFYKKPPATGATVMTGGS-----ATLSKIQ 140
 QY 90 KCDTVRIKGVLGQPTTAPPLMTSEGNVTAEDTEEAIRAFVAVAAASAAEAHWHRLVLL 149
 DB 141 KSSGIPVPVPV-----NGRKTSLDVNSAEFGFLAPGARSNIQ---YRSL--- 181
 QY 150 SQIHEPIGSGGNIINTNGGRSCONPALPSPDQSPGNATSVTRDNYHLTEEEFGVM 209
 DB 182 ----PRPAKSSMSVT---GGRGGPRPVSSSIDPS-----LLSTKQGGILT 219
 QY 210 SQSMKWHSONKSGGVPVRGPTQEPCEISOILKESEFV-----PPTPKENN 255
 DB 220 PSRLKEPTKVASGRTP--APVNTQDREKAKAKAVALDSDNISLKSIGSPSTPKNOA 277
 QY 256 KOEREDENWRLLPPVPV---AETPVSPSVTEIE-----TPLQRIIP--RT 294
 DB 278 SHPTATKLAELPPTPLRTAKSFVKPPSLANLDKVNNSLDLPSSSDTTHASKVPDLHAT 337
 QY 295 ATIAGEPLGCHTFTTSPAFVSHVNLNKRKQLELLREVEWPGRGHMAATCKKL--QVEGQ 352
 DB 338 SSASGGPLPSC-FTSPAPILNINSASFQGLELMSFGSVPKETRMYPKLSGLHRSMESL 396
 QY 353 DRTMSLAAA-----PVREAPPPTCASSEPSVPALPGADPQRSABL 393
 DB 397 QPMNSLPSAFPSSTPV-PTPPAPPAAPTEETEELTWSGSPRAGOL 441

RESULT 14
 AAY85564
 ID AAY85564 standard; Protein; 1837 AA.
 AC AAY85564;
 XX

07-JUL-2000 (first entry)

Human homologue of UNC-53 (Hs-UNC-53/1) sequence.

UNC-53; Caenorhabditis elegans; microtubule; neural regeneration;
 anticancer; anti-neurodegeneration; antifibrotic; anti-adhesive; human;
 antisclerotic; antimetastatic; anti-arthritis; autoimmune disease.

OS Homo sapiens.

Key Location/Qualifiers
 Region 958..1014
 /note= "this region is found to be absent when encoded by
 a variant cDNA isolated from frontal cortex"
 Region 1033..1040
 /note= "this region is found to be absent when encoded by
 a variant cDNA isolated from frontal cortex"
 Region 1173..1175
 /note= "this region is found to be absent when encoded by
 a variant cDNA from Hela or colorectal
 adenocarcinoma tissue"
 Misc-difference 1233
 /label= Leu or Ser

XX WO9963080-A1.
 PN 09-DEC-1999.
 PD 02-JUN-1999; 99WO-EP03848.
 XX 03-JUN-1998; 98GB-0011962.
 XX (JANC) JANSSEN PHARM NV.
 XX Luyten WHML, De Raeymaeker MC, Geysen JJGH, Bogaert TAOE;
 PI Maerten LJS, Verhasselt P, Van De Craen M;
 XX WPI: 2000-116370/10.
 DR N-PSDB; AAA07835.
 XX Novel proteins and nucleic acids e.g. for treating neurodegeneration -
 PS Claim 93; Fig 1b; 146pp; English.
 CC The invention provides vertebrate (human) protein homologue of a UNC-53
 CC protein of Caenorhabditis elegans. The UNC-53 binds to microtubules or
 CC their plus ends. The UNC-53 sequences are used to promote neural
 CC regeneration, revascularization and wound healing; also for treating
 CC neurodegenerative disease, acute traumatic injury, fibrotic disease and
 CC autoimmune diseases (e.g. rheumatoid arthritis and sclerosis). The UNC-53
 CC polynucleotides can be used for recombinant production of the proteins,
 CC as a source of probes for detecting allelic variants and polymorphisms,
 CC for sequencing genomic DNA and for detecting UNC-53 expression; and as
 CC source of therapeutic antisense sequences. Cells that express the
 CC protein are used to identify regulators of cell shape, growth, motility
 CC and migration. They can also be used to identify proteins that are
 CC involved in signal transduction pathways also involving UNC-53, and to
 CC identify compounds that alter attachment of UNC-53 to microtubules. A
 CC target gene coupled to a UNC-53 encoding sequence may be used to deliver
 CC the target gene to a cellular microtubule or its plus ends. The present
 CC sequence represents the amino acid sequence of the first human homologue
 CC of UNC-53, designated hs-UNC-53/1.
 XX
 SQ Sequence 1837 AA;
 Query Match 5.9%; Score 129.5; DB 21; Length 1837;
 Best Local Similarity 21.9%; Pred. No. 0.24;
 Matches 89; Conservative 45; Mismatches 165; Indels 107; Gaps 17;
 QY 41 GAQNLMTQNP-----TLPSVSHRSP---GNAAVSVTGDCCHLTTEEEFGVLVQSM 89
 DB 541 GDRLSDAKKPPSGTARSTSGSGFYKKPPATGATVMTGGS-----ATLSKIQ 591
 QY 90 KCDTVRIKGVLGQPTTAPPLMTSEGNVTAEDTEEAIRAFVAVAAASAAEAHWHRLVLL 149
 DB 592 KSSGIPVPVPV-----NGRKTSLDVNSAEFGFLAPGARSNIQ---YRSL--- 632
 QY 150 SQIHEPIGSGGNIINTNGGRSCONPALPSPDQSPGNATSVTRDNYHLTEEEFGVM 209
 DB 633 ----PRPAKSSMSVT---GGRGGPRPVSSSIDPS-----LLSTKQGGILT 670
 QY 210 SQSMKWHSONKSGGVPVRGPTQEPCEISOILKESEFV-----PPTPKENN 255
 DB 671 PSRLKEPTKVASGRTP--APVNTQDREKAKAKAVALDSDNISLKSIGSPSTPKNOA 728
 QY 256 KOEREDENWRLLPPVPV---AETPVSPSVTEIE-----TPLQRIIP--RT 294
 DB 729 SHPTATKLAELPPTPLRTAKSFVKPPSLANLDKVNNSLDLPSSSDTTHASKVPDLHAT 788
 QY 295 ATIAGEPLGCHTFTTSPAFVSHVNLNKRKQLELLREVEWPGRGHMAATCKKL--QVEGQ 352
 DB 789 SSASGGPLPSC-FTSPAPILNINSASFQGLELMSFGSVPKETRMYPKLSGLHRSMESL 847
 QY 353 DRTMSLAAA-----PVREAPPPTCASSEPSVPALPGADPQRSABL 393
 DB 848 QPMNSLPSAFPSSTPV-PTPPAPPAAPTEETEELTWSGSPRAGOL 892

RESULT 15

RESOLUT
ABR65774

ABB65774 standard: protein: 828 AA.

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AA ABB65774:

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DT 26-MAR-2002 (first entry)

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7) 2007 YILI 07

Prosopbila melanogaster

[illegible]

Drasenhila:

kw
pharmaceutical
biopharmaceutical

pharmaceutical.

XY
30 Dragonfly's molting

US
vv
Drosophila melanogaster

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C - C P V L E L V U C C O R A

PN WO200171042-A2.

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PD 27-SEP-2001.

[illegible]

PF 23~MAR-2001; 2

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PR 23-MAR-2000; 2000US-191637P.

PR 11-JUL-2000; 2000US-0614150.

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PA (PEKE) PE CORP NY.

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PI Venter JC, Adams M, Li PWD, Myers EW;

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DR WPI; 2001-656860/75.

DR N-PSDB; ABL09877.

XX
11/06/2022, 12:55:12 PM

PT New isolated nucleic acid detection reagent for detecting 1000 or more

new isolated nucleic acid detection reagent for detecting 1000 or more genes from *Drosophila* and for elucidating cell signalling and cell-cell

genes from Drosophila and for elucidating cell signaling and cell-cell interactions -

FI INTERACTIONS - XX

PS Disclosure: SF0 TD NO 24114: 2100 + Sequence Listing: English

PS DISCLOSURE; SEQ ID NO Z4114; Z1PP + SEQUENCE LISTING; ENGLISH.
XX

The invention relates to an isolated nucleic acid detection reagent

CC The invention relates to an isolated nucleic acid detection reagent
CC capable of detecting 1000 or more genes from *Proserphila*. The invention is

capable of detecting 1000 or more genes from *Drosophila*. The invention is useful in developmental biology and in elucidating cell signalling and

cell-cell interactions in higher eukaryotes for the development of useful in developmental biology and in elucidating cell signalling and

cell-cell interactions in higher eukaryotes for the development of

CC Insecticides, therapeutics and pharmaceutical drugs. The invention

CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA